

which indicates a low risk

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Date of Report 04-09-2024

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. KOMAL KUMARI					12409030064
Birthday	16-02-2000			Sample ID		11974000
Age at Sample date			24.5	Sample Date		03-09-2024
Gestational age			12+2			
Correction factors						
Fetuses	1 IV	/F		unknown	Previous trisomy 21	unknown
Weight in kg	43 D	iabetes		NO	Pregnancies	unknown
Smoker	NO O	rigin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value	(Corr Mom	Gestational ag	e	12+2
PAPP-A	4.9 m	IU/ml	0.67	Method		LMP
fb-hCG	75.8 ng	g/ml	1.69	Scan date		
Risks at sampling date						
Age Risk			1:971			
Biochemical T21 risk			1:724			
Overall population risk			1:600			
Trisomy 13/18			<1:10000			
Risk				Down's Syndr	ome Risk (Trisomy 21 Sc	reening)
1:10 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT				The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 724 women with the same data, there is one woman with a trisomy 21 pregnancy and 723 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				risk assessment! Calculated risks have no diagnostic values		